## Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application.

- 1. 40. (canceled)
- 41. (previously presented) A pharmaceutical composition comprising a pharmaceutically acceptable excipient or carrier and a compound of Formula I:

$$\begin{array}{c|c} & & & \\ & & & \\ \hline & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & &$$

or a pharmaceutically acceptable salt or prodrug thereof, wherein:

X is O;

Y is CN;

Z is NR<sub>8</sub>R<sub>9</sub>, wherein R<sub>8</sub> and R<sub>9</sub> are independently H or C<sub>1-4</sub> alkyl;

 $R_5$  is hydrogen or  $C_{1-10}$  alkyl;

A is optionally substituted  $C_{6-14}$  aryl; and

B is an optionally substituted indolo ring.

- 42. (previously presented) The pharmaceutical composition of claim 41, wherein A is optionally substituted phenyl.
  - 43. (canceled)

- 44. (original) The pharmaceutical composition of claim 41, wherein X is O, Y is CN and Z is NH<sub>2</sub>.
- 45. (original) The pharmaceutical composition of claim 41, wherein  $R_5$  is hydrogen.
- 46. (previously presented) The pharmaceutical composition of claim 41, comprising a pharmaceutically acceptable excipient or carrier and a compound of Formula II:

$$R_3$$
 $R_4$ 
 $R_5$ 
 $R_2$ 
 $R_1$ 
 $R_1$ 
 $R_2$ 
 $R_1$ 
 $R_2$ 
 $R_3$ 
 $R_4$ 
 $R_5$ 
 $R_5$ 
 $R_5$ 
 $R_5$ 
 $R_5$ 
 $R_7$ 

or a pharmaceutically acceptable salt or prodrug thereof, wherein:

 $R_3$ - $R_4$  are independently hydrogen, halo, haloalkyl, aryl, carbocyclic,  $C_{1-10}$  alkyl, alkenyl, arylalkyl, arylalkenyl, arylalkynyl, carbocycloalkyl, hydroxyalkyl, aminoalkyl, carboxyalkyl, nitro, amino, cyano, acylamido, hydroxy, thiol, acyloxy, azido, alkoxy, carboxy, methylenedioxy, carbonylamido or alkylthiol;  $R_1$  and  $R_2$  taken together with the atoms to which they are attached form a pyrrolo group, wherein said group is optionally substituted;

wherein the aryl portion of said arylalkyl, the aryl portion of said arylalkenyl and the aryl portion of said arylalkynyl are each independently  $C_{6-14}$  aryl; and

said carbocyclic is C<sub>3-8</sub> cycloalkyl or C<sub>3-8</sub> cycloalkenyl.

47. (previously presented) The pharmaceutical composition of claim 46, wherein R<sub>1</sub> and R<sub>2</sub> are taken together to form a structure –CH=CH–N(R)–, wherein R is hydrogen, C<sub>1-10</sub> alkyl, haloalkyl, aryl, fused aryl, carbocyclic, alkenyl, alkynyl, arylalkyl, arylalkenyl, arylalkynyl, carbocycloalkyl, heterocycloalkyl, hydroxyalkyl or aminoalkyl.

## 48.- 49. (canceled)

- 50. (previously presented) The pharmaceutical composition of claim 46, wherein Z is NH<sub>2</sub>.
- 51. (original) The pharmaceutical composition of claim 46, wherein  $R_5$  is hydrogen.
  - 52. (canceled)
- 53. (previously presented) The pharmaceutical composition of claim 46 comprising said compound or a pharmaceutically acceptable salt or prodrug thereof, wherein said optionally substituted  $C_{6-14}$  aryl is

and

- (a) R<sub>10</sub>-R<sub>14</sub> are independently hydrogen, halo, haloalkyl, aryl, fused aryl, carbocyclic, C<sub>1-10</sub> alkyl, alkenyl, alkynyl, arylalkyl, arylalkenyl, arylalkynyl, carbocycloalkyl, hydroxyalkyl, aminoalkyl, carboxyalkyl, nitro, amino, cyano, acylamido, hydroxy, thiol, acyloxy, azido, alkoxy, carboxy, methylenedioxy, carbonylamido or alkylthiol; or
- (b)  $R_{10}$  and  $R_{11}$ , or  $R_{11}$  and  $R_{12}$ , taken together with the atoms to which they are attached form a fused portion of said optionally substituted  $C_{6-14}$  aryl, wherein said fused portion is optionally substituted.
- 54. (previously presented) The pharmaceutical composition of claim 53, wherein  $R_1$  and  $R_2$  are taken together to form a structure –CH=CH–N(R)–, wherein R is hydrogen,  $C_{1-10}$  alkyl, haloalkyl, aryl, fused aryl, carbocyclic, alkenyl, alkynyl, arylalkyl, arylalkynyl, carbocycloalkyl, heterocycloalkyl, hydroxyalkyl or aminoalkyl.

## 55. - 59. (canceled)

60. (previously presented) The pharmaceutical composition of claim 54, wherein  $R_3$ ,  $R_4$  and  $R_5$  are each hydrogen.

61. - 62. (canceled)

- 63. (previously presented) A pharmaceutical composition comprising a pharmaceutically acceptable excipient or carrier and a compound selected from the group consisting of:
- 2-Amino-3-cyano-4-(3-methoxy-4,5-methylenedioxyphenyl)-4*H*-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(2-bromo-4,5-dimethoxyphenyl)-4H-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3-bromo-4,5-dimethoxyphenyl)-4H-indolo[4,5-b]pyran;
- 2-Amino-3-cyano-4-(3-bromo-4,5-dimethoxyphenyl)-8-methyl-4*H*-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3,4,5-trimethoxyphenyl)-4H-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3-nitrophenyl)-4*H*-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3-cyanophenyl)-4H-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3,5-dimethoxyphenyl)-4H-indolo[4,5-b]pyran; and
- 9-Acetamide-2-amino-3-cyano-4-(3-bromo-4,5-dimethoxyphenyl)-4H-indolo[4,5-b]pyran.
  - 64.-65. (canceled).
- 66. (original) The pharmaceutical composition of claim 41, wherein said excipient or carrier is selected from the group consisting of saccharides, starch pastes, gelatin, tragacanth, cellulose preparations, calcium phosphates and polyvinyl pyrrolidone.

- 67. (original) The pharmaceutical composition of claim 66, wherein said excipient or carrier is a saccharide selected from the group consisting of lactose, sucrose, manitol and sorbitol.
- 68. (original) The pharmaceutical composition of claim 41, wherein said excipient or carrier is a lipophilic solvent.
- 69. (original) The pharmaceutical composition of claim 68, wherein said lipophilic solvent is selected from the group consisting of fatty oils, fatty acid esters, polyethylene glycols and paraffin hydrocarbons.
- 70. (original) The pharmaceutical composition of claim 69, wherein said lipophilic solvent is selected from the group consisting of sesame oil, ethyl oleate, triglycerides, polyethylene glycol-400, cremophor and cyclodextrins.
- 71. (original) The pharmaceutical composition of claim 41, wherein said excipient or carrier is selected from the group consisting of vegetable oils, mineral oils, white petrolatum, branched chain fats, branched chain oils, animal fats and high molecular weight alcohol (greater than  $C_{12}$ ).
- 72. (original) The pharmaceutical composition of claim 41, wherein said excipient or carrier is a saline solution.

## 73. - 74. (canceled)

75. (previously presented) A compound of Formula I:

$$\begin{array}{c|c}
A & R_5 \\
\hline
 & X & Z
\end{array}$$
(I)

or a pharmaceutically acceptable salt or prodrug thereof, wherein:

B is optionally substituted indolo;

X is O;

Y is CN;

Z is NR<sub>8</sub>R<sub>9</sub>, wherein R<sub>8</sub> and R<sub>9</sub> are independently H or C<sub>1-4</sub>alkyl;

R<sub>5</sub> is hydrogen or C<sub>1-10</sub> alkyl; and

A is optionally substituted  $C_{6-14}$  aryl.

- 76. (original) The compound of claim 75, wherein said compound is an optionally substituted 4H-indolo[4,5-b]pyran.
- 77. (original) The compound of claim 76, wherein A is optionally substituted phenyl.
- 78. (previously presented) A compound selected from the group consisting of:

- 2-Amino-3-cyano-4-(3-methoxy-4,5-methylenedioxyphenyl)-4*H*-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(2-bromo-4,5-dimethoxyphenyl)-4H-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3-bromo-4,5-dimethoxyphenyl)-4H-indolo[4,5-b]pyran;
- 2-Amino-3-cyano-4-(3-bromo-4,5-dimethoxyphenyl)-8-methyl-4*H*-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3,4,5-trimethoxyphenyl)-4*H*-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3-nitrophenyl)-4*H*-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3-cyanophenyl)-4H-indolo[4,5-b]pyran;
  - 2-Amino-3-cyano-4-(3,5-dimethoxyphenyl)-4H-indolo[4,5-b]pyran; and
- 9-Acetamide-2-amino-3-cyano-4-(3-bromo-4,5-dimethoxyphenyl)-4*H*-indolo[4,5-b]pyran.
  - 79. (canceled)
- 80. (currently amended) The pharmaceutical composition of claim 41, wherein said aryl is selected from the group consisting of phenyl, naphthyl, penanthrenyl phenanthrenyl, anthracenyl, indenyl, azulenyl, biphenyl, biphenylenyl and fluorenyl.
- 81. (curently amended) The compound of claim 75, wherein said aryl is selected from the group consisting of phenyl, naphthyl, penanthrenyl phenanthrenyl, anthracenyl, indenyl, azulenyl, biphenyl, biphenylenyl and fluorenyl.
  - 82. 93. (canceled)

- 94. (previously presented) The pharmaceutical composition of claim 41, wherein said aryl is phenyl.
- 95. (previously presented) The compound of claim 75, wherein said aryl is phenyl.